

# BIONETICS

MUTAGENIC EVALUATION OF COMPOUND FDA 73-83

977051390

MUSTARD, YELLOW

7315 Wisconsin Avenue Bethesda, Maryland 20014

#### LBI PROJECT #2468

### MUTAGENIC EVALUATION OF COMPOUND FDA 73-83

977051390

MUSTARD, YELLOW

#### SUBMITTED TO

FOOD AND DRUG ADMINISTRATION
DEPARTMENT OF HEALTH, EDUCATION AND WELFARE
ROCKVILLE, MARYLAND

SUBMITTED BY

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MAY 30, 1975



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DATE: May 30, 1975

SPONSOR: Food and Drug Administration, Contract Number 223-74-2104

SUBJECT: Evaluation of Test Compound 977051390, Mustard, Yellow FDA 73-83

#### I. OBJECTIVE

The objective of this study was to evaluate the test compound for genetic activity in microbial assays with and without the addition of mammalian metabolic activation preparations.

#### II. MATERIALS

A. Test Compound

Date Received: August, 1974

2. Description: yellow powder

B. <u>Indicator Microorganisms</u>

The following strains of indicator microorganisms were used in the evaluation:

Yeast Strain: Saccharomyces cerevisiae, strain D4

Bacteria Strains: Salmonella typhimurium, strains: TA-1535

TA-1537 TA-1538

### C. Reaction Mixture

The following reaction mixture was employed in the activation tests:

<u>Component</u>	Final Concentration/ml
<ol> <li>TPN (sodium salt)</li> <li>Isocitric acid</li> <li>Tris buffer, pH 7.4</li> <li>MgCl<sub>2</sub></li> </ol>	6.0 μM 49.0 μM 28.0 μM 1.7 μM
5. Tissue homogenate fraction	72.0



#### D. Tissue Homogenates and Supernatants

The tissue homogenates and 9,000 x  $\underline{g}$  supernatants were prepared from tissues of the following mammalian species: Mouse-ICR random bred adult males; rat-Sprague-Dawley adult males; and primate-Macaca mulatta adult males.

#### E. Positive Control Compounds

Table 1 lists chemicals for positive controls in the direct and activation assays.

# TABLE 1 POSITIVE CONTROLS USED IN DIRECT AND ACTIVATION ASSAYS

Assay	<u>Chemical<sup>a</sup></u>	Solvent	Probable Mutagenic Specificity
Nonactivation	Ethyl methanesulfonate 2-Nitrofluorene Quinacrine mustard	Water or saline Dimethylsulfoxide <sup>C</sup> Water or saline	BPS <sup>b</sup> FS <sup>b</sup>
Activation	Dimethylnitrosamine 2-Acetylaminofluorene	Water or saline Dimethylsulfoxide <sup>C</sup>	BPS <sup>b</sup> · FS <sup>b</sup>

a Concentrations given in the Results Section

Previously shown to be non-mutagenic

#### III. METHODS

#### A. <u>Toxicity</u>

The solubility, toxicity and doses for all chemicals were determined prior to screening.

Each chemical was tested for survival against the specific indicator strains over a range of doses to determine the 50% survival dose. Bacteria were tested in phosphate buffer, pH 7.4, for one hour at 37°C on a shaker. Yeasts were tested in phosphate buffer, pH 7.4, for four hours at 30°C on a shaker. The 50% survival curve and the 1/4 and 1/2 50% doses calculated.

If no toxicity was obtained for a chemical with a given strain, then a maximum dose of 5% (w/v) was used against the strain.

Unless otherwise specified, the doses calculated for the tests in buffer were applied to the activation tests. The solubility of the test chemical under treatment conditions is stated in the Results Section.



BPS = base-pair substitution; FS = frameshift

#### B. Plate Tests

In the nonactivation procedure, approximately 10° cells of a log-phase culture of the bacterial indicator strains were spread over the surface of a minimal plate, and a measured amount of the test chemical was placed in the center of the test plate. In activation tests, the test chemical was added to the cells, and an aliquot of the mixture was spread on the surface of the test plate. The reaction mixture (0.1 ml) plus tissue extract was then spotted on the surface of the plate. Positive and solvent controls were included. All plates were incubated at 37°C for four days and then scored. Each compound (test, positive control and solvent control) was done in duplicate. Concentrations of the positive control compounds are listed in the Results Section.

#### C. <u>Suspension Tests</u>

#### 1. Nonactivation

Log-phase bacteria and stationary-phase yeast cultures of the indicator organisms were grown in complete broth, washed and resuspended in 0.9% saline to densities of 1 x  $10^9$  cells/ml and 5 x  $10^7$  cells/ml, respectively. This constituted the working stock for tests of a group of test chemicals and their respective controls. Tests were conducted in plastic tissue culture plates. Cells plus appropriate volume(s) of the test chemical were added to the wells to give a final volume of 1.5 ml. The solvent replaced the test chemical in the negative controls. was at 30°C for four hours for yeast tests and at 37°C for one hour for bacterial tests. All flasks were shaken during treatment. Following treatment, the plates were set on ice. Aliquots of cells were removed, diluted in sterile saline ( $4^{\circ}$ C) and plated on the appropriate complete media. Undiluted samples from flasks containing the bacteria were plated on minimal selective medium in reversion experiments. Samples from a 10<sup>-1</sup> dilution of treated cells were plated on the selected media for enumeration of gene conversion with strain D4. Bacterial plates were scored after incubation for 48 hours at 37°C. The yeast plates were incubated at 30°C for 3-5 days before scoring.

#### 2. Activation

Bacteria and yeast cells were grown and prepared as described in the nonactivation tests. Measured amounts of the test and control chemicals plus 0.25 ml of the stock-cell suspension were added to wells of the Linbro plate containing the appropriate tissue fraction and reaction mixture. All flasks (bacteria and yeast) were incubated at 37°C in an oxygen atmosphere with shaking. The treatment times as well as the dilutions, plating procedures and scoring of the plates were the same as described for nonactivation tests.



### D. <u>Preparation of Tissue Homogenates and 9,000 x g Cell Fractions</u>

Male animals (sufficient to provide the necessary quantities of tissues) were killed by cranial blow, decapitated and bled. Organs were immediately dissected from the animal using aseptic techniques and placed in ice-cold 0.25 M sucrose buffered with Tris at pH of 7.4. Upon collection of the desired quantity of organs, they were washed twice with fresh buffered sucrose and completely homogenized with a motor-driven homogenizing unit at 4°C. The whole organ homogenate obtained from this step was divided into two samples. One sample was frozen at -80°C and the other was centrifuged for 20 minutes at 9,000 x g in a refrigerated centrifuge. The supernatant from the centrifuged sample was retained and frozen at -80°C. These two frozen samples were used for the activation studies.

#### E. <u>Data Recording and Reporting</u>

Following the specified incubation periods all population plates were scored by an automatic colony counter and the results from each plate of a set were recorded, in ink, on data processing forms. All minimal or other types of selective media plates were hand scored and the results recorded along with the respective population data. Other relevant experimental data were recorded on experimental definition forms. For bacteria strains the number of colonies recorded from either the population or selective plates represents that number in 1 ml of test suspension plated. The numbers recorded for the yeast strain D4 represent the number in 0.5 ml of test suspension plated. The data were then processed and printed from a computer program.



- IV. RESULTS SECTION
- A. <u>Solubility Properties of the Test Compound</u>
- 1. Name or code designation of the test compound: 977051390, Mustard
  - Yellow
- 2. Test solvent: DMSO
- 3. Solubility of the test compound under treatment conditions: Chemical soluble at treatment concentrations.
- 4. Additional comments: yellow powder
- B. <u>Toxicity and Dosage Determinations for the Test Compound</u>
- 1. Test date for toxicity determination: February 11, 1975
- 2. The 50% survival level was determined for bacteria and yeast indicator organisms by conducting survival curves with the test compound at the following concentrations:

#### Percent Concentration (w/v or v/v)

5.0

0.5

0.05

0.005

0.0005

3. Concentrations of the test compound used in the mutagenicity tests:

•	Percent Concentration			
<u>Dose</u>	Bacteria	Yeast		
1/4 50% Survival	0.25	0.0025		
1/2 50% Survival	0.50	0.0050		
50% Survival	1.00	0.0100		
Plate Tests	0.50	•		



#### SUMMARY OF TEST RESULTS ٧.

### Plate Tests

Name or code designation of the test compound: 977051390 A.

Test date: March 29, 1975 В.

Concentration of the test compound: C. 0.5%

• •		J. J	, c copot	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	0.36	-			
Test		<u>Species</u>	<u>Tissue</u>	TA.	<u>-1535</u>	TA	<u>-1537</u>	ŢA	<u>-1538</u>
1.	Nonactivation			<u> </u>	2	1	<u>2</u>	1	<u>2</u>
	Solvent Control Positive Control <sup>a</sup> Test Compound			11 >10 <sup>5</sup> 6	20 >10 <sup>5</sup> 13	4 193 8	5 207 8	15 145 17	11 134 16
2.	Activation								
	Negative Control Solvent Control Reaction Mixture			7 20	10 24	6 5	3 9	7 15	12 14
	Control			23	20	7	8	18	16
	Positive Control Positive Control Positive Control	Mouse	Liver Lung Testes	>10 <sup>3</sup> 12 16	>10 <sup>3</sup> 8 17	39 2 3	34 1 8	343 11 7	357 16 11
	Positive Control Positive Control Positive Control	Rat	Liver Lung Testes	>10 <sup>3</sup> 14 16	>10 <sup>3</sup> 7 13	89 2 5	88 3 7	347 14 10	341 18 12
	Positive Control Positive Control Positive Control	Monkey	Liver Lung Testes	273 8 15	356 8 12	30 2 3	33 2 6	123 13 8	119 14 11
	Test Compound Test Compound Test Compound	Mouse	Liver Lung Testes	10 3 8	8 10 7	2 0 2	4 1 1	9 14 41	17 9 17
	Test Compound Test Compound Test Compound	Rat	Liver Lung Testes	11 4 10	8 10 8	3 1 4	3 2 1	10 15 13	15 7 17
	Test Compound Test Compound Test Compound	Monkey	Liver Lung Testes	9 3 5	7 10 4	2 0 3	4 2 3	13 16 8	16 8 15
a	TA-1537 QM 20	μl/plate μg/plate μg/plate	TA	N-1535 N-1537 N-1538	DMNA AAF AAF	100	μM/plate μg/plate μg/plate	<u> </u>	



### DATA TABLE TERMS AND ABBREVIATIONS

OR TERM	DEFINITION OR EXPLANATION
COMPOUND	Client designated compound number appears in this column.
TEST CODES	NAN = Nonactivation: Solvent Control NAP = Nonactivation: Positive Control NA1 = Nonactivation: Test Compound Dose 1 NA2, etc. = Reflects the other dose level(s)
	A+C = Negative Chemical Control A-C = Activation: Solvent Control ACP = Activation: Positive Control ACT = Activation: Test Compound A+T = Activation: Tissue Control
	LI = Liver Tissue Activation Fraction  LU = Lung Tissue Activation Fraction  KI = Kidney Tissue Activation Fraction  TE = Testes Tissue Activation Fraction  1,2, etc. = Dose Levels
CONCENTRATION	All test compound dose levels are expressed as a whole number followed by an exponent (negative) identified by the appropriate units.
	Example: 0025-2PCT = 0.25 percent concentration
POPU	Total number of viable cells in the plating sample raised to some exponent printed directly below the abbreviation (i.e., EP + $6 = x \cdot 10^6$ ).
MUT 1	Total number of mutants or convertants obtained from the sample plated raised to some exponent printed directly below the abbreviation (i.e., EP + 0 = $\times$ 100). For strain D4, MUT 1 represents th number of ADE+ convertants.
MUT 2	Only used for strain D4 and represents the number of TRY+ convertants in the plated sample.
FREQ 1	The calculated mutation or gene conversion frequency times the negative exponent written directly below. For strain D4, FREQ 1 represents the ADE+ value.
FREQ 2	Only used for strain D4 and represents the TRY+ conversion frequency.
CONTAM	Presence of contamination on any plates.

### cDATA TABLE TERMS AND ABBREVIATIONS (continued)

OR TERM	DEFINITION OR EXPLANATION
AAF	2-Acetylaminofluorene
DMSO	Dimethylsulfoxide
DMN	Dimethylnitrosamine
EMS	Ethyl Methanesulfonate
QМ	Quinacrine Mustard
NF	Nitrofluorene
SPECIES	Animal Strains
SPRDAW	Sprague Dawley Rats

Flow ICR Random Bred Mice

New Zealand White Rabbit

Dog, Mixed Breed

Rhesus Monkey ( $\underline{Macaca}$   $\underline{mulatta}$ )



**ICRFLO** 

**RHESUS** 

MIXEDB

NEWZEA

## LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM REPORT EXR34

COMPOUND FREQUENCY SUMMARY REPORT 05/15/75

SPECIES

COMPOUND 977051390

TEST	ORG	TA1535 HIS EX-8	TA1537 HIS EX-8	TA1538 HIS EX-8	0000D4 ADE EX-5	0000D4 TRY EX-5
NAN		10.15	2.10	9.02	1.58	1.69
NAP		273.70	225.89	145.97	113.25	60.11
NA1		17.37	2.87	15.72	1.00	5.26
NA2		14.99	2.04	11.66	1.32	3.07



### LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM REPORT EXR34

COMPOUND FREQUENCY SUMMARY REPORT 05/15/75

SPECIES ICRFLO COMPOUND 977051390

TEST	DRG	TA1535 HIS EX-8	TA1537 HIS EX-8	TA1538 HIS EX-8	0000D4 ADE EX-5	0000D4 TRY EX-5	•
ACT	A+C	5.99	2.10	11.20	16.01	4.11	
ACT	A+T	12.45	2.10	10.46	15.24	3.26	
ACT	A-C	6.65	2.91	10.54	19.85	3.33	•
ACT	PL I	1168.86	10.07	37.43	20.62	4.92	
ACT	PLU	8.06	2.19	12.57	39.08	3.22	
ACT	PTE	9.32	2.86	12.54	23.53	3.29	
ACT	LII	4.95	2.75	27.16	17.96	2.25	
ACT	LI2	7.18	2.26	25.07	19.01	3.09	
ACT	LU1	6.52	2.22	22.13	16.37	1.79	
ACT	LU2	8.86	2.29	19.48	20.21	1.38	,
ACT	TE1	6.07	4.24	29.11	13.80	2.51	
ACT	TE2	9.20	2.03	21.79	15.50	2.88	



## LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM REPORT EXR34

COMPOUND FREQUENCY SUMMARY REPORT 06/24/75

SPECIES SPRDAW/RAT

\*COMPOUND 977051390

TEST	ORG	TA1538 HIS EX-8	TA1535 HIS EX-8	TA1537 HIS EX-8	0000D4 ADE EX-5	0000D4 TRY EX-5
ACT	A+C	5.39	17.47	0.63	0.58	0.39
ACT	A+T	10.55	11.03	1.23	2.35	2.75
ACT	A-C	5.36	11.62	0.58	0.72	0.18
ACT	PL I	59.37	1500.00	7.28	8.75	10.22
ACT	PLU	7.31	18.29	0.84	3.40	1.39
ACT	PTE	8.13	22.47	0.33	3.38	1.93
ACT	LI1	6.74	11.87	0.50	1.47	3.19
ACT	LI2	10.92	11.02	0.59	2.81	5.88
ACT	LU1	5.75	14.66	0.48	1.88	2.26
ACT	LU2	5.44	12.32	0.64	1.13	1.25
ACT	TE 1	7.36	15.92	0.62	2 • 48	0.78
ACT	TE2	6.70	13.88	0.82	2.39	1.74



## LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM REPORT EXR34

COMPOUND FREQUENCY SUMMARY REPORT 05/15/75

SPECIES RHESUS

COMPOUND 977051390

TEST	ORG	TA1535 HIS EX-8	TA1537 HIS EX-8	TA1538 HIS EX-8	0000D4 ADE EX-5	0000D4 TRY EX-5
ACT	A+C	7.43	7.44	4.49	10.40	1.65
ACT	A+T	14.05	11.84	4.09	15.69	2.55
ACT	A-C	7.01	7.04	5.55	14.76	2.24
ACT	PLI	1352.79	19.31	24.13	14.99	2.71
ACT	PLU	4.62	6.35	2.77	12.58	2.35
ACT	PTE	8.16	8.79	6.29	20.07	2.79
ACT	LI1	4.50	9.13	2.92	20.00	3.33
AÇT	LI2	6.88	3.94	3.97	18.33	2.05
ACT	LU1	7.47	19.41	4.04	16.61	2.68
ACT	. LU2	7.30	10.74	1.99	13.30	2.75
ACT	TE 1	8.29	7.49	6.61	16.96	3.39
ACT	TE2	7.31	5.54	14.18	18.55	2.73



#### VI. INTERPRETATION OF RESULTS AND CONCLUSIONS

Compound FDA 73-83, Mustard Yellow, was evaluated for genetic activity in a series of in vitro microbial assays with and without metabolic activation. The following results were obtained:

- Α. Salmonella typhimurium
- 1. Plate Tests

At a concentration of 0.5%, this compound was not mutagenic for any of the indicator strains in direct or activation plate tests.

2. Nonactivation suspension tests

The results of these tests were negative.

Activation suspension tests 3.

The results of these tests were considered negative. The mutant frequencies for several of the doses with TA-1538 using mouse tissues were slightly increased. The rat and primate data were clearly negative. No reason was apparent for the increases.

- В. Saccharomyces cerevisiae
- 1. Nonactivation suspension tests

The results of these tests were negative.

2. Activation suspension tests

The results of these tests were negative. Random apparent increases were noted at a few of the ade doses. No pattern was observed indicating that the increases were induced.

C. Conclusions

The test compound, Mustard Yellow, did not exhibit genetic activity in the in vitro assays employed in this evaluation.

Submitted by:

David Brusick, Ph.D. Director of Genetics

### APPENDIX

Tabulation of Data





CONTRACT 22374-2104					PROJECT 02468				
EXPERIMENT 505803		DETECTOR TA1535 SPECIES		DATE - 05/15/75					
		ORG		POPU	MUT1	FREQ1			
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8	CONTAM		
•	NAN		SALINE	1340	0136	10.15	0		
	NAP		EMS 0.002 %	1327	3632	273.70	0		
977051390	NA1		0005-1 PCT.	0547	0095	17.37	0		
977051390	NA2		0025-2 PCT.	0647	0097	14.99	0		



EXPERI	COMENT 508		22374-2104 DETECTOR TA1537 SPECIES			PROJECŤ 02468 DATE - 05/15/75		
_		ORG		POPU	MUT1	FREQ1	03/13/13	
COMPOU	ND TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8	CONTAM	
	NAN		SALINE	0713	0015	2.10	0	
	NAP		OM 1.0 UG/ML	0421	0951	225.89	0	
977051	390 NA1		0005-1 PCT.	0767	0022	2.87	1	
977051	390 NA2		0025-2 PCT.	1029	0021	2.04	0	



			22374-2104	PROJECT 02468					
EXPERIMEN	T 5058	02	DETECTOR TA1538	SPE	CIES	DATE -	05/15/75		
		ORG		POPU	MUT1	FREQ1			
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8	CONTAM		
	NAN		DMSO	0599	0054	, 9•02	0		
	NAP		NF 125 UG-ML	0546	0797	145.97	0		
977051390	NA1		0005-1 PCT.	0229	0036	15.72	0		
977051390	NA2		0025-2 PCT.	0386	0045	11.66	0		



EXPERIMEN			22374-2104 DETECTOR 0000D4	PROJECT 02468 SPECIES DATE - 05/15/75						
COMPOUND		ORG ID	CONCENTRATION	POPU EP+4	MUT1 EP+1	MUT2 EP+1	FREQ1 EP-5	FREQ2 EP-5	CONTAM	
	NAN		SALINE	0947	0015	0016	1.58	1.69	0	
	NAP		EMS 1.0 %	0702	0795	0422	113.25	60.11	7	
977051390	NAI		0005-3 PCT.	0399	0004	0021	1.00	5.26	1	
977051390	NA2		0025-4 PCT.	0456	0006	0014	1.32	3.07	1	



CONTRAC EXPERIMENT 506301			22374-2104 DETECTOR TA1535	PROJECT 02468 SPECIES ICRFLO DATE - 05/15/75					
		ORG		POPU	MUT1	FREQ1			
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8	CONTAM		
	A+C		DMN 50 UM/ML	0734	0044	5.99	0		
	A+T	•	***NO MATCH***	0819	0102	12.45	0		
	A-C		SALINE	0917	0061	6.65	o		
	ACP	LI	DMN 50 UM/ML	0456	5330	1168.86	1		
	ACP	LU	DMN 50 UM/ML	0608	0049	8.06	0		
·	ACP	TE	DMN 50 UM/ML	0590	0055	9•32	0		
977051390	ACT	LII	0005-1 PCT.	0808	0040	4.95	0		
977051390	ACT	LI2	0025-2 PCT.	0850	0061	7.18	0		
977051390	ACT	LU1	0005-1 PCT.	0951	0062	6.52	2		
977051390	ACT	LU2	0025-2 PCT.	0677	0060	8.86	· O		
977051390	ACT	TE1	0005-1 PCT.	0791	0048	6.07	0		
977051390	ACT	TE2	0025-2 PCT.	0815	0075	9.20	0		



			22374-2104			PROJECT 02468	
EXPERIMENT	5059	01	DETECTOR TA1537	SPE	CIES ICR	FLO DATE	- 05/15/75
		ORG		POPU	MUT1	FREQ1	
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8	CONTAM
	A+C		AAF 800 UG/ML	1476	0031	2.10	0
	A+T		***NO MATCH***	1524	0032	2.10	• 0
	A-C		DMSO	1445	0042	2.91	0
	ACP	LI	AAF 800 UG/ML	1489	0150	10.07	0
	ACP	LU	AAF 800 UG/ML	1463	0032	2.19	0
٠	ACP	TE	AAF 800 UG/ML	1293	0037	2.86	0
977051390	ACT	LII	0005-1 PCT.	1493	,0041	2.75	2
977051390	ACT	LI2	0025-2 PCT.	1546	0035	2.26	2
977051390	ACT	LU1	0005-1 PCT.	1530	0034	2.22	2
977051390	ACT	LU2	0025-2 PCT.	1483	0034	2.29	2
977051390	ACT	TE1	0005-1 PCT.	1485	0063	4.24	o
977051390	ACT	TE2	0025-2 PCT.	1476	0030	2.03	0



EXPERIMEN			22374-2104 DETECTOR TA1538	PROJECT 02468 SPECIES ICRFLO DATE - 05/15/75					
COMPOUND	TEST	ORG ID	CONCENTRATION	POPU EP+6	MUT1 EP+0	FREO1 EP-8	CONTAM		
	A+C		AAF 800 UG/ML	0598	0067	11.20	0		
	A+T		***NO MATCH***	0841	0088	10.46	0		
	A-C		DMSO	0446	0047	10.54	0		
	ACP	LI	AAF 800 UG/ML	0505	0189	37.43	0		
	ACP	LU	AAF 800 UG/ML	0565	0071	12.57	2		
	ACP	TE	AAF 800 UG/ML	0606	0076	12.54	0		
977051390	ACT	LI1	0005-1 PCT.	0243	0066	27.16	2		
977051390	ACT	LI2	0025-2 PCT.	0335	0084	25.07	0		
977051390	ACT	LU1	0005-1 PCT.	0235	0052	22.13	2		
977051390	ACT.	LU2	0025-2 PCT.	0267	0052	19.48	2		
977051390	ACT	TE1	0005-1 PCT.	0213	0062	29.11	0		
977051390	ACT	TE2	0025-2 PCT.	0234	0051	21.79	2		



	CON	ITRACT	22374-2104			PRO.	JECT 0246	58	
EXPERIMEN.	T 5104	01	DETECTOR 0000D4	. SPE	CIES I	CRFLO	DA	ATE - 05/	15/75
COMPOUND	TEST	ORG ID	CONCENTRATION	POPU	MUT1	MUT2	FREQ1	FREQ2	
COMPOUND	1631	10	CONCENTRATION	EP+4	EP+1	EP+1	EP-5	EP-5	CONTAM
	A+C		DMN 90 UM/ML	0706	0113	0029	16.01	4.11	0
	A+T		***NO MATCH***	1319	0201	0043	15.24	3.26	. 6
	A-C		SALINE	0660	0131	0022	19.85	3.33	2
·	ACP	LI	DMN 90 UM/ML	0325	0067	0016	20.62	4.92	6
	ACP	LU	DMN 90 UM/ML	0870	0340	0028	39.08	3.22	2
	ACP	TE	DMN 90 UM/ML	0850	0200	0028	23.53	3.29	4
977051390	ACT	LII	0005-3 PCT.	0579	0104	0013	17.96	2.25	0
977051390	AC T	LI2	0025-4 PCT.	0647	0123	0020	19.01	3.09	6
977051390	ACT	LU1	0005-3 PCT.	0391	0064	0007	16.37	1.79	0
977051390	ACT	LU2	0025-4 PCT.	0579	0117	0008	20.21	1.38	0
977051390	ACT	TE1	0005-3 PCT.	0558	,0077	0014	13.80	2.51	4
977051390	ACT	TE2	0025-4 PCT.	0729	0113	0021	15.50	2.88	6



EXPERIMENT			22374-2104 DETECTOR TA1535	SPE	CIES SPR	PROJECT 02468 RDAW DATE - (	05/15/75
		ORG		POPU	MUTI	FREQ1	
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8	, CONTAM
•	A+C		DMN 50 UM/ML	0435	0076	17.47	0
·	A+T		***NO MATCH***	0571	0063	11.03	0
	A-C		SALINE	0697	0081	11.62	0
	ACP	LI	DMN 50 UM/ML	0388	5820	1500.00	0.
	ACP	LU	DMN 50 UM/ML	0410	0075	18.29	0
	ACP	TE	DMN 50 UM/ML	0356	0080	22.47	2
977051390	ACT	LI1	0005-1 PCT.	0851	0101	11.87	2
977051390	ACT	LI2	0025-2 PCT.	0708	0078	11.02	2 .
977051390	ACT	LU1	0005-1 PCT.	0682	0100	14.66	2
977051390	ACT	LU2	0025-2 PCT.	0641	0079	12.32	2
977051390	ACT	TE 1	0005-1 PCT.	0622	0099	15.92	2
977051390	ACT	TE2	0025-2 PCT.	0706	0098	13.88	2



		22374-2104	PROJECT 02468					
EXPERIMENT	5070	01	DETECTOR TA1537	SPE	CIES SPRDAW	DATE	- 05/15/75	
COMPOUND	7507	DRG		POPU	MUT1	FREQ1		
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8	CONTAM	
	A+C		AAF 800 UG/ML	1906	0012	0.63	0	
	A+T		***NO MATCH***	2198	0027	1.23	0	
	A-C		DMSO	1900	0011	0.58	o	
	ACP	LI	AAF 800 UG/ML	1594	0116	7.28	0	
	ACP	LU	AAF 800 UG/ML	1657	0014	0.84	2	
	ACP	TE	AAF 800 UG/ML	1841	0006	0.33	2	
977051390	ACT	LII	0005-1 PCT.	1989	0010	0.50	o	
977051390	ACT	LI2	0025-2 PCT.	2219	0013	0.59	o	
977051390	ACT	LU1	0005-1 PCT.	2063	0010	0.48	0	
977051390	ACT	LU2	0025-2 PCT.	1881	0012	0.64	o	
977051390	ACT	TEl	0005-1 PCT.	2084	0013	0.62	0	
977051390	ACT	TE2	0025-2 PCT.	1699	0014	0.82	1	



EXPERIMENT			22374-2104 DETECTOR TA1538	SPE	CIES SPR	PROJECT 02468 DAW DATE - 0	5/15/75
COMPOUND	TEST	ORG ID	CONCENTRATION	POPU EP+6	MUT1 EP+0	FREQ1 EP-8	CONTAM
	A+C		AAF 800 UG/ML	0445	0024	5.39	0
	A+T		***NO MATCH***	0275	0029	10.55	2
	A-C		DMSO	0392	0021	5.36	0
	ACP	LI	AAF 800 UG/ML	0507	0301	59•37	0
	ACP	LU	AAF 800 UG/ML	0588	0043	7.31	0
	ACP	TE	AAF 800 UG/ML	0541	0044	8.13	0
977051390	ACT	LII	0005-1 PCT.	0430	0029	6.74	0
977051390	ACT	LI2	0025-2 PCT.	0293	0032	10.92	0
977051390	ACT	LU1	0005-1 PCT.	0435	0025	5.75	0
977051390	ACT	LU2	0025-2 PCT.	0570	0031	5.44	2
977051390	AC T	TE1	0005-1 PCT.	0421	0031	7.36	2
977051390	ACT	TE2	0025-2 PCT.	0522	0035	6.70	0



<b></b>			22374-2104				JECT 024	68		
EXPERIMENT	5066	01	DETECTOR 0000D4	SPE	CIES S	PRDAW/	RAT		DATE - 06/24/75	,
COMPOUND	TEST	ORG	CONCENTRATION	POPU	MUT1	MUT2	FREQ1	FREQ2		
COMPOUND	1531	ID	CONCENTRATION	EP+4	EP+1	EP+1	EP-5	EP-5	CONTAM	
	A+C		DMN 90 UM/ML	1037	0006	0004	0.58	0.39	4	
	A+T		***NO MATCH***	1020	0024	0028	2.35	2.75	4	
	A-C		SALINE .	1104	8000	0002	0.72	0.18	4	
	ACP	L I	DMN 90 UM/ML	0949	0083	0097	8.75	10.22	4	
	ACP	LU	DMN 90 UM/ML	0794	0027	0011	3.40	1.39	6	
	ACP	TE	DMN 90 UM/ML	1035	0035	0020	3.38	1.93	4	
977051390	ACT	LI1	0005-3 PCT.	0814	0012	0026	1.47	3.19	4	
977051390	ACT	LI2	0025-4 PCT.	0782	0022	0046	2.81	5.88	4	
977051390	ACT	LU1	0005-3 PCT.	0797	0015	0018	1.88	2.26	4	
977051390	ACT	LU2	0025-4 PCT.	0798	0009	0010	1.13	1.25	, <b>4</b>	
977051390	ACT	TE1	0005-3 PCT.	0766	0019	0006	2.48	0.78	4	
977051390	ACT	TE2	0025-4 PCT.	0922	0022	0016	2.39	1.74	o	



·	CON	ITRACT	22374-2104	PROJECT 02468				
EXPERIMENT	5072	201	DETECTOR TA1535	. SPE	CIES RHE	SUS DATE -	05/15/75	
		ORG		POPU	MUT1	FREQ1		
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8	CONTAM	
	A+C		DMN 50 UM/ML	0888	0066	7 • 43	2	
	A+T		***NO MATCH***	0541	0076	14.05	0	
	A-C		SALINE	0913	0064	7.01	0	
	ACP	LI	DMN 50 UM/ML	0502	6791	1352.79	0	
	ACP	LU	DMN 50 UM/ML	0823	0038	4.62	0	
	ACP	TE	DMN 50 UM/ML	0625	0051	8.16	0	
977051390	ACT	LII	0005-1 PCT.	0666	0030	4.50	0	
977051390	ACT	LI2	0025-2 PCT.	0828	0057	6.88	2	
977051390	ACT	LU1	0005-1 PCT.	0495	0037	7.47	0	
977051390	ACT	LU2	0025-2 PCT.	0671	0049	7.30	2	
977051390	ACT	TE1	0005-1 PCT.	0543	0045	8.29	0	
977051390	ACT	TE2	0025-2 PCT.	0916	0067	7.31	o	



. 6	EXPERIMENT	-		22374-2104 DETECTOR TA1537.	SPE		ECT 02468 DATE - 05/15/	75
			ORG		POPU	MUT1	FRE01	
(	COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8 (	MATNO
		A+C		AAF 800 UG/ML	0847	0063	7.44	2
		A+T		***NO MATCH***	0515	0061	11.84	0
		A-C		DMSO	0710	0050	7.04	0
		ACP	LI	AAF 800 UG/ML	0844	0163	19.31	0
		ACP	LU	AAF 800 UG/ML	0914	0058	6.35	0
		ACP	TE	AAF 800 UG/ML	0842	0074	8.79	0
9	77051390	ACT	LII	0005-3 PCT.	0449	0041	9.13	0
ç	977051390	ACT	LI2	0025-4 PCT.	1421	0056	3.94	0
9	77051390	ACT	LU1	0005-3 PCT.	0407	0079	19.41	0
ç	977051390	ACT	LU2	0025-4 PCT.	0447	0048	10.74	0
9	77051390	ACT	TEL	0005-3 PCT.	0574	0043	7.49	0
ç	77051390	ACT	TE2	0025-4 PCT.	0686	0038	5.54	0



EXPERIMENT		NTRACT 501	22374-2104 DETECTOR TA1538	SPE	CIES RHES	PROJECT 02468 SUS DATE -	05/15/75
		ORG		POPU	MUT1	FREQ1	
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8	CONTAM
	A+C		AAF 800 UG/ML	0802	0036	4.49	<b>O</b>
	A+T		***NO MATCH***	0464	0019	4.09	0
	A-C		DMSD	0721	0040	5.55	0
	ACP	LI	AAF 800 UG/ML	0601	0145	24.13	2
	ACP	LU	AAF 800 UG/ML	0649	0018	2.77	0
	ACP	TE	AAF 800 UG/ML	0572	0036	6.29	0
977051390	ACT	LI1	0005-1 PCT.	0513	0015	2.92	2
977051390	ACT	LIZ	0025-2 PCT.	0453	0018	3.97	. 0
977051390	ACT	LU1	0005-1 PCT.	0421	0017	4.04	0
977051390	ACT	LU2	0025-2 PCT.	0654	0013	1.99	0
977051390	ACT	TEl	0005-1 PCT.	0439	0029	6.61	O
977051390	ACT	TE2	0025-2 PCT.	0395	0056	14.18	0



CONTRACT			22374-2104 PROJECT 02468							
EXPERIMEN	T 5105	01	DETECTO	R 0000D4	SPE	CIES R	RHESUS	D	ATE - 05/	15/75
		ORG			POPU	MUT1	MUT2	FREQI	FREQ2	
COMPOUND	TEST	ID	CONCENT	RATION	EP+4	EP+1	EP+1	EP-5	EP-5	· CONTAM
	A+C		DMN 90 I	UM/ML	0423	0044	.0007	10.40	1.65	o
,	A+T		***NO M	ATCH***	0548	0086	0014	15.69	2.55	0
	A-C		SALINE		0759	0112	0017	14.76	2.24	0
	ACP	LI	DMN 90 1	UM/ML	0774	0116	0021	14.99	2.71	0
	ACP	LU	DMN 90 (	JM/ML	0469	0059	0011	12.58	2.35	0
	ACP	TE	DMN 90 (	JM/ML	0538	0108	0015	20.07	2.79	0
977051390	ACT	LII	0005-3	PCT.	0570	0114	0019	20.00	3.33	4
977051390	ACT	LI2	0025-4	PCT.	0633	0116	0013	18.33	2.05	0
977051390	ACT	LUI	0005-3 F	PCT.	0560	0093	0015.	16.61	2.68	0
977051390	ACT	LU2	0025-4 F	PCT.	0436	0058	0012	13.30	2.75	0
977051390	ACT	TE1	0005-3 F	PCT.	0560	0095	0019	16.96	3.39	0
977051390	ACT	TE2	0025-4 F	PCT.	0550	0102	0015	18.55	2.73	0 .